FORM PTO-1449 U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

INFORMATION DISCLOSURE STATEMENT BY APPLICANT

(Use several sheets if necessary)

ATTY. DOCKET NO.:	APPLICATION NO.:
OC01617K	10/654,546
APPLICANT: Timothy J. Guzi *et al.*	
FILING DATE: 09/03/2003	GROUP: 1624

U.S. PATENT DOCUMENTS

*EXAMINER INITIAL		DOCUMENT NUMBER	DATE	NAME	CLASS	SUB-CLASS	FILING DATE IF APPROPRIATE
	AA	US 5,571,813	11/05/1996	Rühter *et al.*			
	AB	US 5,602,136	02/11/1997	Rühter *et al.*			
	AC	US 5,602,137	02/11/1997	Rühter *et al.*			
	AD	US 5,688,949	11/18/1997	Inoue *et al.*			
	AE	US 5,707,997	01/13/1998	Shoji *et al.*			
	AF	US 5,919,815	07/06/1999	Bradley *et al.*			
	AG	US 6,040,321	03/21/2000	Kim *et al.*			
	AH	US 6,191,131	02/20/2001	He *et al.*			
	AI	US 6,262,096	07/17/2001	Kim *et al.*			
	AJ	US 6,413,974	07/02/2002	Dumont *et al.*			

FOREIGN PATENT DOCUMENTS

		DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUB-CLASS	TRANSLATION YES	TRANSLATION NO
	AK	EP 0 628 559	04/03/2002	Europe				
	AL	EP 1 334 973	08/13/2003	Europe				
	AM	WO 02/40485	05/23/2002	PCT				
	AN	WO 02/50079	06/27/2002	PCT				
	AO	WO 95/35298	12/28/1995	PCT				

OTHER DOCUMENTS *(Including Author, Title, Date, Pertinent Pages, Etc.)*

	AP	Vesely *et al.*, "Inhibition of Cyclin-Dependent Kinases by Purine Analogues", *Eur. J. Biochem* (1994), **224**: 771-786.
	AQ	Kim *et al.*, "Discovery of Aminothiazole Inhibitors of Cyclin-Dependent Kinase 2: Synthesis, X-ray Crystallographic Analysis, and Biological Activities", *Journal of Medical Chemistry*, Page EST:22.3, A-W.
	AR	Mettey *et al.*, "Aloisines, a New Family of CDK/GSK-3 Inhibitors. SAR Study, Crystal Structure in Complex with CDK2, Enzyme Selectivity, and Cellular Effects", *J. Med. Chem.* (2003), **46**(2): 222-236.
	AS	Novinson *et al.*, "Synthesis and Antifungal Properties of Certain 7-Alkylaminopyrazolo[1,5-α]pyrimidines", *J. Med. Chem.* (1977), **20**(2): 296-299.
	AT	Senderowicz *et al.*, "Phase I Trial of Continuous Infusion Flavopiridol, a Novel Cyclin-Dependent Kinase Inhitibor, in Patients with Refractory Neoplasms", *Journal of Clinical Oncology* (September 1998), **16**(9): 2986-2999.
	AU	Meijer *et al.*, "Biochemical and Cellular Effects of Roscovitine, a Potent and Selective Inhibitor of the Cyclin-Dependent Kinases CDC2, CDK2 and CDK5", *Eur. J. Biochem.* (1997), **243**:527-536.
	AV	Bible *et al.*, "Cytotoxic Synergy Between Flavopiridol (NSC 649890, L86-8275) and Various Antineoplastic Agents: The Importance of Sequence of Administration", *Cancer Research* (August 15, 1997), **57**: 3375-3380.
	AW	Shiota *et al.*, "Synthesis and Structure-Activity Relationship of a New Series of Potent Angiotensin II Receptor Antagonists: Pyrazolo[1,5-α]pyrimidine Derivatives", *Chem. Pharm. Bull.* (1999), **47**(7): 928-938.
	AX	Yasuo Makisumi, "Studies on the Azaindolizine Compounds. XI. Synthesis of 6,7-Disubstituted Pyrazolo[1,5-α]pyrimidines.", *Chem. Pharm. Bull* (1962), **10**: 620-626.

EXAMINER	DATE CONSIDERED
	6/17/05

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

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U.S. PATENT DOCUMENTS

*EXAMINER INITIAL		DOCUMENT NUMBER	DATE	NAME	CLASS	SUB-CLASS	FILING DATE IF APPROPRIATE
	AA	6,107,305	08/22/2000	Misra *et al.*			
	AB	6,413,974 B1	07/02/2002	Dumont *et al.*			
	AC						
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	AH						
	AI						
	AJ						
	AK						

FOREIGN PATENT DOCUMENTS

		DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUB-CLASS	TRANSLATION YES	TRANSLATION NO
		DE 102 23 917 A1	12/11/2003	Germany				
	AL	EP 0 628 559 A	12/14/1994	Europe				
	AM	WO 02/50079 A1	06/27/2002	WIPO				
	AN	WO 03/091256 A1	11/06/2003	WIPO				
	AO							
	AP							

OTHER DOCUMENTS *(Including Author, Title, Date, Pertinent Pages, Etc.)*

	AQ	Translation of WO 03/91256, *A Rising Sun Communications Ltd. Translation Product*, (1-62)
	AR	
	AS	
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	AU	
	AV	

EXAMINER	DATE CONSIDERED
/Jon McElwain/	6/17/07

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